Amendment to the Claims

- 1. (*Cancelled*) A catheter comprising:
- (a) an elongate catheter body having a distal end and a proximal end, the elongate catheter body including an indicator lumen and an insulating lumen, the indicator lumen having a restricted cross sectional area adjacent the distal end of the elongate catheter body;
 - (b) a dilution sensor connected to the elongate catheter body; and
- (c) a guide wire extending through the restricted cross sectional area of the indicator lumen to project from the distal end of the elongate catheter body.
- 2. (*Cancelled*) The catheter of Claim 1, wherein the dilution sensor is a thermistor.
- 3. (*Cancelled*) The catheter of Claim 1, wherein the insulating lumen is at least partially intermediate the indicator lumen and the dilution sensor.
- 4. (*Cancelled*) The catheter of Claim 1, wherein the elongate catheter body is configured as a retrograde catheter.
- 5. (Currently amended) A catheter <u>assembly</u> comprising:
- (a) an elongate catheter body having a distal end and a proximal end, the elongate catheter body including an indicator lumen <u>having a terminal port and a radial injection port intermediate the terminal port and the proximal end of the catheter body</u>; [[and]]

- (b) a guide wire extending through a length of the indicator lumen; and
- (c) a controller connected to one of the catheter body and the guide wire, the controller selected to compensate for passage of an indicator from the indicator lumen through the terminal port.
- 6. (Currently amended) The catheter <u>assembly</u> of Claim 5, wherein the indicator lumen terminates at the distal end of the elongate catheter body.
- 7. (Currently amended) The catheter <u>assembly</u> of Claim 5, wherein the indicator lumen has a reduced cross sectional area adjacent the distal end of the elongate catheter body.
- 8. (Currently amended) The catheter <u>assembly</u> of Claim 5, wherein the indicator lumen has a reduced cross sectional area adjacent the distal end of the elongate catheter body and the guide wire is sized to be slideably received through the reduced cross sectional area.
- 9. (Currently amended) The catheter <u>assembly</u> of Claim 5, wherein the indicator lumen has a reduced cross sectional area adjacent the distal end of the elongate catheter body and the guide wire is sized to reduce passage of [[an]] <u>the</u> indicator through the reduced cross sectional area of the indicator lumen.
- 10. (Currently amended) The catheter <u>assembly</u> of Claim 5, wherein the indicator lumen terminates adjacent the distal end of the elongate catheter body.

- 11. (*Cancelled*) The catheter of Claim 5, wherein the indicator lumen includes a terminal port at the distal end of the elongate catheter body and a radial injection port spaced from the terminal port.
- 12. (Currently amended) The catheter <u>assembly</u> of Claim 5, further comprising a dilution sensor connected to the elongate catheter body.
- 13. (Currently amended) The catheter <u>assembly</u> of Claim 12, wherein the dilution sensor is a thermistor.
- 14. (Currently amended) A method of introducing an indicator through a catheter, the method comprising:
- (a) passing a guide wire through an indicator lumen in an elongate catheter body to pass a portion of the guide wire through a terminal port of the indicator lumen; [[and]]
- (b) passing the indicator through the indicator lumen to pass from the elongate catheter body through the terminal port and an injection port intermediate the terminal port and a proximal end of the catheter body; and
 - (c) compensating for passage of the indicator through the terminal port.
- 15. (*Cancelled*) The method of Claim 14, further comprising simultaneously passing the guide wire and the indicator through the indicator lumen.
- 16. (Currently amended) The method of Claim 14, further comprising passing the guide wire through a reduced cross sectional area of the indicator lumen-to reduce passage of the indicator there through.

- 17. (Currently amended) The method of Claim 14, further comprising <u>passing</u> the indicator through the indicator lumen to contact a portion of <u>simultaneously locating</u> the guide wire and the indicator in the indicator lumen.
- 18. (Currently amended) The method of Claim 14, further comprising passing the guide wire through a reduced cross sectional area of the indicator lumen to increase a flow of the indicator through the a radial injection port.
- 19. (New) The method of Claim 14, wherein compensating for passage of the indicator through terminal port includes compensating for a volume of the indicator passing through the terminal port.
- 20. (New) The method of Claim 14, wherein compensating for passage of the indicator through terminal port includes compensating for a volume of the indicator passing through the terminal port corresponding to the relationship $Q = \frac{k(T_b T_i) \cdot V(1-a)}{S}, \text{ where } Q \text{ is a blood flow rate, } k \text{ is a coefficient related to}$ thermal capacity of a measured flow and the indicator, T_b is the temperature of the measured flow prior to injection, T_i is the temperature of the indicator prior to entering the measured flow, V is the volume of the indicator, S is the area under the temperature versus time curve resulting from the mixing of the indicator and a is the portion of the indicator passing through the terminal port.

- 21. (New) The method of Claim 14, wherein compensating for passage of the indicator through terminal port includes compensating for a thermal effect of the indicator passing through the terminal port.
- 22. (New) The method of Claim 14, wherein compensating for passage of the indicator through terminal port includes compensating for a thermal effect of the indicator passing through the terminal port corresponding to the relationship $Q = \frac{k(T_b T_i) \cdot V(1 a)}{(S_m S_{in})}$, where Q is a blood flow rate, k is a coefficient related to thermal capacity of a measured flow and the indicator, T_b is the temperature of the measured flow prior to injection, T_i is the temperature of the indicator prior to entering the measured flow, V is the volume of the indicator, S_m is the total area under the temperature versus time curve resulting from the mixing of the indicator, S_{in} is the part of the area under the dilution curve related to a cooling thermal change of a sensor inside the catheter body and a is the portion of the indicator passing through the terminal port.
- 23. (New) The catheter assembly of Claim 5, wherein a portion of the guide wire is in the terminal port.
- 24. (New) The catheter assembly of Claim 5, wherein the catheter body further includes a spacer lumen intermediate the distal end and the proximal end.

- 25. (New) The catheter assembly of Claim 5, wherein a portion of the guide wire is disposed in the terminal port and the controller is selected to compensate for passage of the indicator through the terminal port.
- 26. (New) The catheter assembly of Claim 5, wherein the controller is selected to compensate for passage of a volume of the indicator through the terminal port corresponding to the relationship $\mathcal{Q} = \frac{k(T_b, T_i) \cdot V(1-a)}{S}$, where Q is a blood flow rate, k is a coefficient related to thermal capacity of a measured flow and the indicator, T_b is the temperature of the measured flow prior to injection, T_i is the temperature of the indicator prior to entering the measured flow, V is the volume of the indicator, S is the area under the temperature versus time curve resulting from the mixing of the indicator and a is the portion of the indicator passing through the terminal port.
- 27. (New) The catheter assembly of Claim 5, wherein the controller is selected to compensate for passage of a volume of the indicator through the terminal port corresponding to the relationship $Q = \frac{k(T_{b-}T_i)\cdot V(1-a)}{(S_m-S_{in})}$, where Q is a blood flow rate, k is a coefficient related to thermal capacity of a measured flow and the indicator, T_b is the temperature of the measured flow prior to injection, T_i is the temperature of the indicator prior to entering the measured flow, V is the volume of the indicator, S_m is the total area under the temperature versus time curve resulting from the mixing of the indicator, S_{in} is the part of the area

under the dilution curve related to a cooling of a sensor inside the catheter body and a is the portion of the indicator passing through the terminal port.